

**HYDROCORTISONE ACETATE- hydrocortisone acetate suppository**  
**AARNA USA INC.**

*Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, click here.*

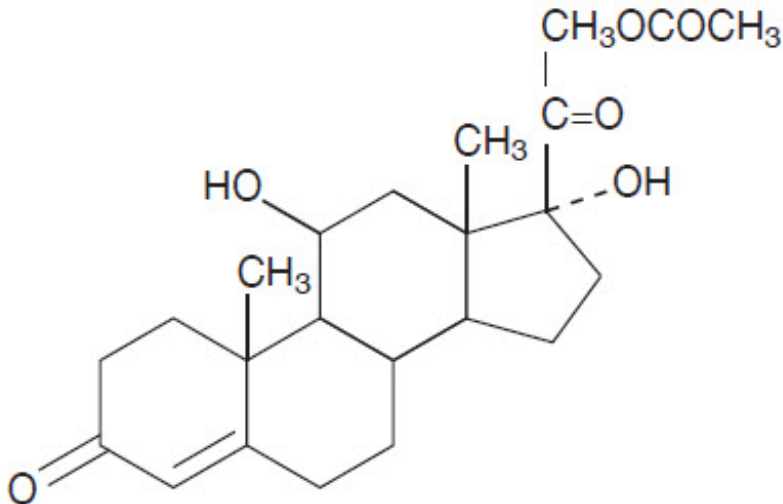
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**Hydrocortisone Acetate Suppositories 25mg and 30mg**

**For Rectal Administration**

**Rx Only**

**DESCRIPTION**

Hydrocortisone acetate is a corticosteroid designed chemically as pregn-4-ene-3, 20-dione, 21-(acetyloxy)-11, 17-dihydroxy-(11 $\beta$ ) with the following structural formula:



Each 25 mg suppository for rectal administration contains hydrocortisone acetate, USP 25 mg in a specially blended hydrogenated vegetable base.

Each 30 mg suppository for rectal administration contains hydrocortisone acetate, USP 30 mg in a specially blended hydrogenated vegetable base.

**CLINICAL PHARMACOLOGY**

In normal subjects, about 26% of hydrocortisone acetate is absorbed when the suppository is applied to the rectum. Absorption of hydrocortisone acetate may vary across abraded or inflamed surfaces.

Topical steroids are primarily effective because of their anti-inflammatory, anti-pruritic and vasoconstrictive action.

**INDICATIONS AND USAGE**

Hydrocortisone acetate suppositories are indicated for use in inflamed hemorrhoids, post-irradiation (factitial) proctitis, as an adjunct in the treatment of chronic ulcerative colitis, cryptitis, other inflammatory conditions of anorectum, and pruritus ani.

## **CONTRAINDICATIONS**

Hydrocortisone acetate suppositories are contraindicated in those patients having a history of hypersensitivity to hydrocortisone acetate or any of the components.

## **PRECAUTIONS**

Do not use hydrocortisone acetate suppositories unless adequate proctologic examination is made.

If irritation develops, the product should be discontinued and appropriate therapy instituted.

In the presence of an infection, the use of an appropriate antifungal or antibacterial agent should be instituted. If a favorable response does not occur promptly, hydrocortisone acetate should be discontinued until the infection has been adequately controlled.

## **Carcinogenesis**

No long-term studies in animals have been performed to evaluate the carcinogenic potential of corticosteroid suppositories.

## **Pregnancy Category C**

In laboratory animals, topical steroids have been associated with an increase in the incidence of fetal abnormalities when gestating females have been exposed to rather low dosage levels. There are no adequate and well-controlled studies in pregnant women.

Hydrocortisone acetate suppositories should only be used during pregnancy if the potential benefit justifies the risk of the fetus. Drugs of this class should not be used extensively on pregnant patients, in large amounts, or for prolonged periods of time.

It is not known whether this drug is excreted in human milk, and because many drugs are excreted in human milk and because of the potential for serious adverse reactions in nursing infants from hydrocortisone acetate suppositories, a decision should be made whether to discontinue nursing or to discontinue the drug, taking into account the importance of the drug to the mother.

## **ADVERSE REACTIONS**

The following local adverse reactions have been reported with hydrocortisone acetate suppositories; burning, itching, irritation, dryness, folliculitis, hypopigmentation, allergic contact dermatitis, secondary infection.

To report an adverse event, please contact AARNA USA INC at 1-877-225-6999.

## **DRUG ABUSE AND DEPENDENCE**

Drug abuse and dependence have not been reported in patients treated with hydrocortisone acetate suppositories.

## **OVERDOSAGE**

If signs and symptoms of systemic overdose occur, discontinue use.

## **DOSAGE AND ADMINISTRATION**

### **FOR RECTAL ADMINISTRATION**

Detach one suppository from strip of suppositories. Hold suppository upright and carefully separate tabs at top opening and pull downward from the pointed end to expose the suppository. Remove the suppository from the pocket. Avoid excessive handling of suppository which is designed to melt at body temperature. Insert one suppository rectally, pointed end first. Insert one suppository in the rectum twice daily, morning and night for two weeks, in nonspecific proctitis. In more severe cases, one suppository three times a day or two suppositories twice daily. In factitial proctitis, the recommended duration of therapy is six to eight weeks or less, according to the response of the individual case.

## **HOW SUPPLIED**

Hydrocortisone acetate suppositories 25mg are off-white, smooth surfaced and bullet shaped with one pointed end.

Box of 12 suppositories, NDC 82568-0150-1

Box of 24 suppositories, NDC 82568-0150-2

Hydrocortisone acetate suppositories 30mg are off-white, smooth surfaced and bullet shaped with one pointed end.

Box of 12 suppositories, NDC 82568-0160-1

## **STORAGE**

Store at 20°-25°C (68°-77°F) [See USP Controlled Room Temperature]. Excursions permitted to 15°- 30°C (59°-86°F). Store away from heat. Protect from freezing. Avoid contact with eyes.

## **KEEP THIS AND ALL DRUGS OUT OF THE REACH OF CHILDREN**

In case of accidental ingestion, seek professional assistance or contact a Poison Control Center immediately.

**PHARMACIST:** This product is not an Orange Book rated product, therefore all prescriptions using this product shall be subject to state and federal statutes as applicable. This product has not been subjected to FDA therapeutic or other equivalency testing. There are no claims of bioequivalence or therapeutic equivalence. Each person recommending a prescription substitution using this product shall make such

recommendation based on his/her professional knowledge and opinion, upon evaluating the active ingredients, inactive ingredients, excipients and chemical information contained within the enclosed prescribing information.

Rx Only

**Manufactured for:**  
AARNA USA INC.

Rev. 04/26

## PRINCIPAL DISPLAY PANEL - 25 mg Suppository Carton

NDC 82568-0150-1

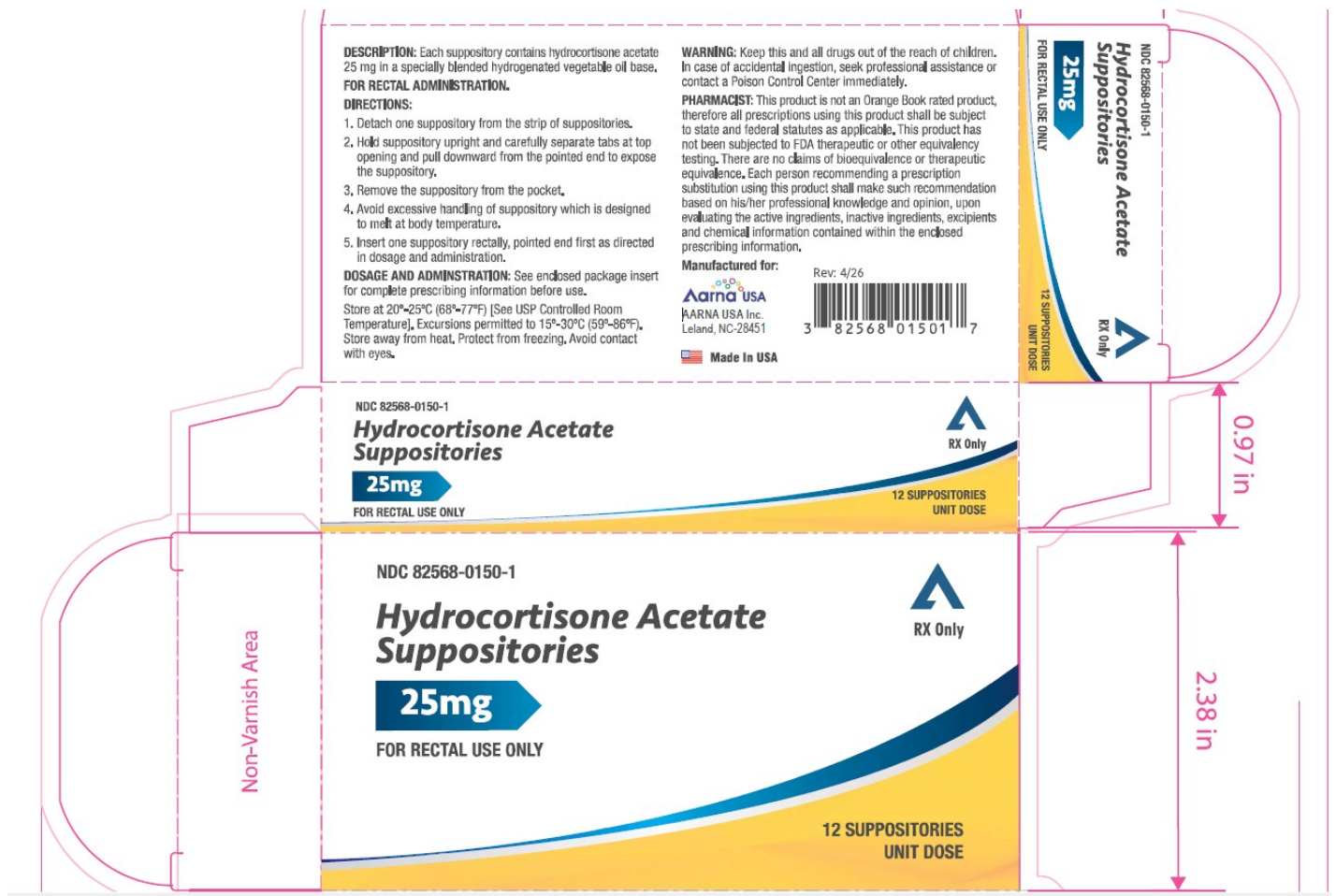
Rx Only

Hydrocortisone Acetate  
Suppositories 25 mg

FOR RECTAL USE ONLY

Aarna USA Inc

12 Suppositories  
Unit Dose



NDC 82568-0150-2

Rx Only

Hydrocortisone Acetate  
Suppositories 25 mg

FOR RECTAL USE ONLY

Aarna USA Inc.

24 Suppositories  
Unit Dose



**PRINCIPAL DISPLAY PANEL - 30 mg Suppository Carton**

NDC 82568-0160-1

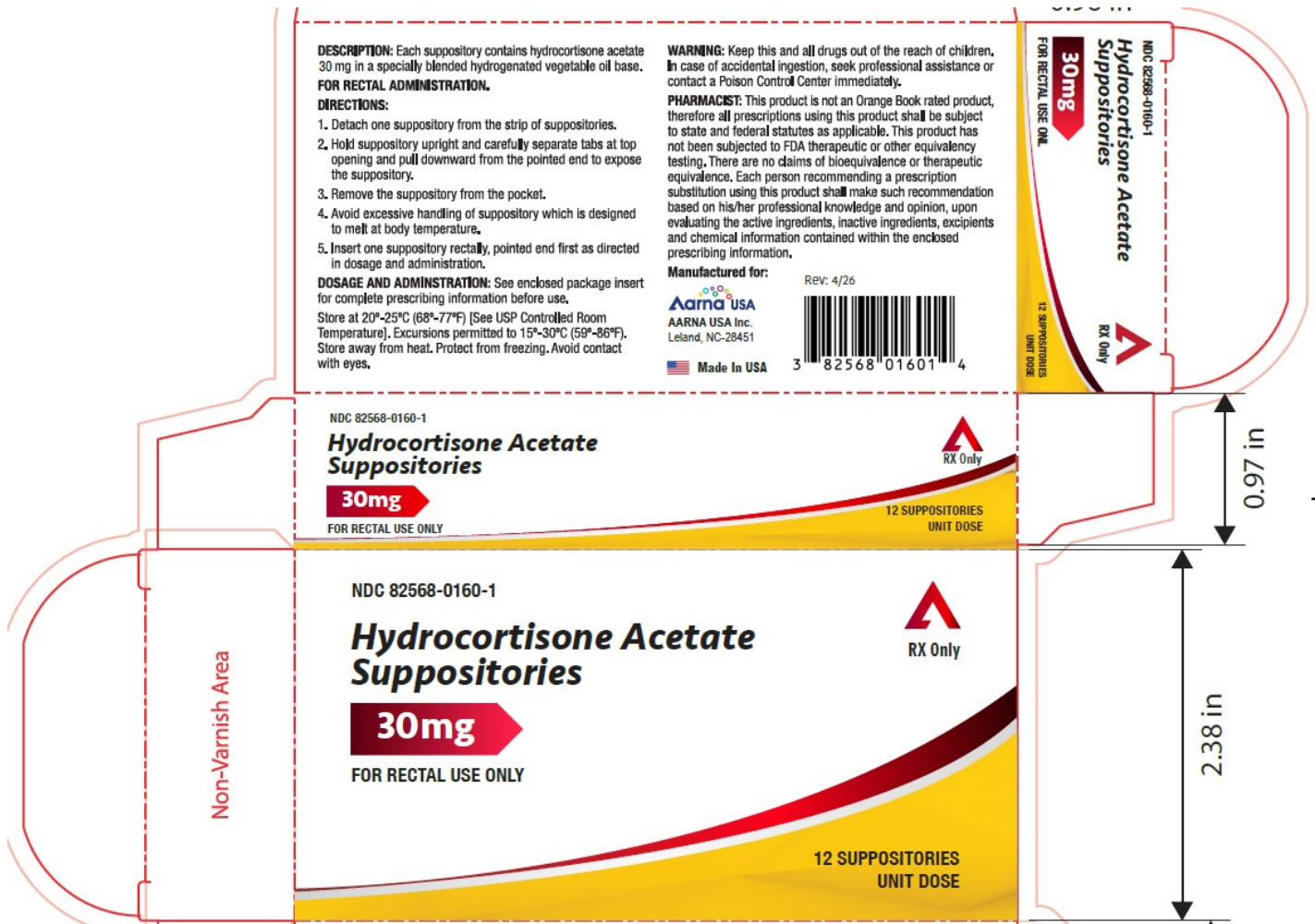
Rx Only

Hydrocortisone Acetate  
Suppositories 30 mg

FOR RECTAL USE ONLY

Aarna USA Inc

12 Suppositories  
Unit Dose



## HYDROCORTISONE ACETATE

hydrocortisone acetate suppository

### Product Information

<b>Product Type</b>	HUMAN PRESCRIPTION DRUG	<b>Item Code (Source)</b>	NDC:82568-0150
<b>Route of Administration</b>	RECTAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
HYDROCORTISONE ACETATE (UNII: 3X7931PO74) (HYDROCORTISONE - UNII:W4X0X7BPJ)	HYDROCORTISONE ACETATE	25 mg

### Inactive Ingredients

Ingredient Name	Strength
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**HYDROGENATED PALM KERNEL OIL** (UNII: FM8D1RE2VP)

### Product Characteristics

<b>Color</b>	white (Off-White)	<b>Score</b>	
<b>Shape</b>	BULLET	<b>Size</b>	
<b>Flavor</b>		<b>Imprint Code</b>	
<b>Contains</b>			

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:82568-0150-1	12 in 1 CARTON; Type 0: Not a Combination Product	04/16/2026	
2	NDC:82568-0150-2	24 in 1 CARTON; Type 0: Not a Combination Product	04/16/2026	

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		04/16/2026	

## HYDROCORTISONE ACETATE

hydrocortisone acetate suppository

### Product Information

<b>Product Type</b>	HUMAN PRESCRIPTION DRUG	<b>Item Code (Source)</b>	NDC:82568-0160
<b>Route of Administration</b>	RECTAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>HYDROCORTISONE ACETATE</b> (UNII: 3X7931PO74) (HYDROCORTISONE - UNII:W4X0X7BPJ)	HYDROCORTISONE ACETATE	30 mg

### Inactive Ingredients

Ingredient Name	Strength
<b>HYDROGENATED PALM KERNEL OIL</b> (UNII: FM8D1RE2VP)	

### Product Characteristics

<b>Color</b>	white (Off-White)	<b>Score</b>	
<b>Shape</b>	BULLET	<b>Size</b>	

<b>Flavor</b>		<b>Imprint Code</b>	
<b>Contains</b>			

**Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:82568-0160-1	12 in 1 CARTON; Type 0: Not a Combination Product	04/16/2026	

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Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		04/16/2026	

**Labeler - AARNA USA INC. (118515992)**

Revised: 4/2026

AARNA USA INC.